SOROKIN, M.F.; KHINCHINA, E.L.

Synthesis of allyl glycidyl ester and butyl methacrylate copclymers. Lakokras. mat. i ith prim. no.4:5-10 '63. (MIRA 16:10)

SOROKIN, M.F.; KOCHNOV, I.M.

Synthesis of glycidyl methacrylate and styrene copolymers in solvents and their use as a base for the manufacture of protective coatings. Lakokras. mat. i ikh prim. no.4:10-15 '63.

SOROKIN, M.F.; MIKHAYLOVA, L.S.

Polymerization of aryl glycidol esters in the presence of metal halides. Lakokras. mat. i ikh prim. no.5:10-15 '63. (MIRA 16:11)

L 9839-63 EPR/EWP(j)/EPF(c)/EWT(m)/BDS-AFFTC/ASD-Pg-li/Pc-li/Pr-li-ACCESSION NR: AP3000395 S/0191/63/000/005/0011/0014

AUTHOR: Sorokin, M. F.; Latov, V. K.; Korkishko, Zh. T.; Kochnova, Z. A.

TITLE: Copolymers of unsaturated ethers of glycidol. Copolymerization of methyl methacrylate with allylglycidyl ethers in solutions

SOURCE: Plasticheskiye massy\*, no. 5, 1963, 11-14

TOPIC TAGS: copolymerization, methyl methacrylate, 2-progenyl 2,3-epoxypropyl, copolymerization rate, reaction temperature, initiator concentration, reactivity ratios, yields, molecular weight, benzoyl peroxide

ABSTRACT: Methyl methacrylate (MMA) and 2-propenyl 2,3-epoxyypropyl ether (PEPE) have been copolymerized in methyl ethyl ketone, dioxane, or toluene solution at 70 to 90C in the presence of 0.5 to 1.0 mol% of benzoyl peroxide or Alpha, Alpha-azobisisobutyronitrile. The reaction was conducted under nitrogen in solutions whose initial concentration of the monomers was 30%, with MMA and PEPE in ratios of 2:1, 1:1, and 1:2. The copolymerization rate dropped with an increase in the PEPE content and increased with an increase

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L 9839-63 ACCESSION NR: AP3000395

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in the reaction temperature or initiator concentration. Because MMA is much more reactive than PEPE, the MMA-PEPE ratio in the copolymers is much higher than that in the initial monomer mixture. The monomer reactivity ratios were calculated to be 40.7 for MMA and 0.035 for PEPE. The copolymers, obtained in yields of 41 to 86, are white solids readily soluble in benzene, toluene, acetone, or dioxane. Their molecular weights vary from 2000 to 10,000, decreasing with an increase in the initial PEPE concentration, reaction temperature, or initiator concentration. Copolymers prepared in dioxane solution have the highest molecular weight. Orig. art. has: 5 figures and 4 tables.

ASSOCIATION: none

SUBMITTED: 00

DATE ACQ: 10Jun63

ENCL: 00

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NO REF SOV: 000

OTHER: Oll

ار عدر/ja Card 2/2

8/0303/64/000/002/0007/0010

ACCESSION NR: AP4034711

AUTHORS: Sorokin, M. F.; Kochnova, Z. A.; Korkishko, Zh. T.

TITLE: Two-component polyurethane lacquers on the base of glycidyl ester polymers, synthesized in the presence of two- and three-functional initiators

SOURCE: Lakokrasochny\*ye materialy\* i ikh primeneniye, no. 2, 1964, 7-10

TOPIC TAGS: polyurethane lacquer, glycidyl ester, phenylglycidyl ester, butylglycidyl ester, glycidyl ester polymer, diisocyanate, triisocyanate, chemical film stability, lacquer film hardening, lacquer film adhesion

ABSTRACT: The base n-butylglycidyl ester polymer (BGEP) and the phenylglycidyl ester polymer (PGEP) were synthesized from the corresponding monomers by block polymerization at 90C in the presence of the initiators resorting, trimethylolpropane, and NaOH. While 30% resorcinol yielded linear polymers of an average molecular weight of 570, the same amount of trimethylolpropane produced branched polymers of molecular weights within the 760-775 range. The BCEP and PCEP polymers so produced were used to make lacquers by being blended with toluilenediisocyanate (TD), with 4,4'-diphenylmethanediisocyanate (DMD), or with 4,4',4"triphenylmethanediisocyanate (TMD). Since the hardening of films of such two-

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ACCESSION NR: AP4034711

component compositions is caused by the formation of urethanes, the authors performed a series of tests, using various NCO:OH ratios, temperatures, and reaction periods. It was found that at 90C satisfactorily hardened films with high physicomechanical properties and good appearance were obtained by blending the glycidyl esters of linear structure with TMD, and by blending the polymers of branched structure with TD or DMD. It was also found that for blends of the linear PGE polymer with TMD the optimum ratio of NCO:OH was 1.3:1. This yielded (at 90C) a completely hardened film within 6-7 hours. For the branched BGE and PGE polymers, the optimal ratios of NCO:OH in their blends with DMD were 1.4:1 and 1.1:1, and the hardening periods were 1 and 3 hours respectively. A period of 16-20 hours was hardening periods were 1 and 3 hours respectively. A period of 16-20 hours was required for hardening films of the branched BGE polymer blended with TD at a required for hardening films of the branched BGE polymer blended with TD at a required for hardening films of the branched BGE polymer blended with TD at a required for hardening films of the branched BGE polymer blended with TD at a required for hardening films of the branched BGE polymer blended with TD at a required for hardening films of the branched BGE polymer blended with TD at a required for hardening films of the branched BGE polymer blended with TD at a required for hardening films of the branched BGE polymer blended with TD at a required for hardening films of the branched BGE polymer blended with TD at a required for hardening films of the branched BGE polymer blended with TD at a required for hardening films of the branched BGE, and 3 formulas.

ASSOCIATION: none SUBMITTED: CO SUB CODE: MT

DATE ACQ: 20May64 NO REF 80V: 004 ENCL: 00

Card 2/2

ACCESSION NR: AP4040513

5/0303/64/000/003/0008/0010

AUTHOR: Zlobina, V. R.; Sorokin, M. F.; Blagonravova, A. A.

TITLE: Curing of epoxide resins

SOURCE: Lakokrasochny\*ye materialy\* i ikh primeneniye, no. 3, 1964, 8-10

TOPIC TAGS: epoxide resin, phenyl glycidyl ether, carbanilide, epoxy group, imino group epoxide resin curing

ABSTRACT: It was established that phenyl glycidyl ether reacts with carbanilide to form hydroxyl-containing compounds as a result of the interaction between epoxy and imino groups; it was postulated that in addition, a reaction is possible between the epoxy groups and the hydroxyl groups formed. It was shown that condensation of butyl ether takes place in the course of the reaction of phenyl glycidyl ether with dibutyl ester of dimethylolurea, simultaneously with the interaction of the epoxy and imino groups, and with the interaction of the hydroxyl groups formed and the epoxy groups. This condensation is indicated by the evolution of butanol in the course of the reaction.

ASSOCIATION: none

Card 1/2

ACCESSION NR: AP4043819

\$/0303/64/000/004/0001/0004

AUTHOR: Sorokin, M. F.; Kochnova, Z. A.; Korkishko, Zh. T.; Vasil'yeva, O. A.

TITLE: Premixed polyurethan coatings based on polymers of n-butyl- and phenylglycidyl ethers.

FOURCE: Lakokrasochny\*ye materialy\* i ikh primeneniye, no. 4, 1964. 1-4

TOPIC TAGS: premixed coating, prepolymer synthesis, cured film, chemically resistant coating, polymerized phenylglycidyl ether, polymerized n-butyl ether, toluylene diisocyanate, polyurethan, polyurethan film

ABSTRACT: Two types of prepolymers with isocyanate end groups were synthesized from linear or branched polymers of n-butyl or phenylglycidyl ethers and toluylene diisocyanate, using 30 mol. % trimethylolpropane or phenylglycidyl or resorcinol as initiators, respectively, to obtain premixed polyurethan coatings characterized by stability in storage. Two type-1 prepolymers were synthesized from the branched polymers at 60°C. Both the polymer and the diisocyanate were used as 50% solutions in xylene, the temperature of the exothermic reaction did not exceed 20°C, and the reaction was complete 1.5 hrs. after toluylene diisocyanate was added. Two type-2 prepolymers were synthesized at 70°C from

Card 1/2

SOROKIN, M.F.; MIKHAYLOVA, L.S.

Polymerization of phenyl glycidyl ether in the presence of aluminum isopropylate and zinc chloride. Vysokom.sced. 6 no.4:677-683 Ap 164. (MIRA 17:6)

Moskovskiy krimiko∴tekhnologicheskiy institut imeni
 D.I. Mendeleyeva.

L 16038-65 EWT(m)/EWP(j)/T Pc-4 RM S/0303/64/000/005/0001/0005

AUTHOR: Sorokin, M. F., Korkishko, Zh. T., Kochnova, Z.A.

TITLE: Two-component polyurethan varnishes based on nitrogenous tetrafunctional polymers of glycidyl ethers and polyisocyanates

SOURCE: Lakokrasochny\*ye materialy\* i ikh primeneniye, no. 5, 1964, 1-5

TOPIC TAGS: lacquer, polyurethane varnish, nitrogenous polymer, tetrafunctional polymer, glycidyl ether, polyisocyanate, tolylene disocyanate, hexamethylene disocyanate, hexamethylene diamine, ethylene diamine, tolylene diamine, polymer hardening, polymerization initiator

ABSTRACT: Tetrafunctional nitrogen-containing polymers based on glycidyl ethers were synthesized and were shown to be of possible use for obtaining two-component polyurethan compositions with tolylene and hexamethylene disocyanate. The polymers were synthesized in the presence of tetrafunctional initiators such as hexamethylene diamine, ethylene diamine and m-tolylene diamine. It was found that the addition of a small amount of sodium hydroxide (5 mol. %) accelerates the process at a molar ratio of 7:1 of ether to diamine. The structural formulas of the polymers are given. The characteristics of the polymers and copolymers synthesized in the presence of diamines and a catalyst (5 mol. % NaOH) are

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ACCESSION NR: AP4047671

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tabulated. Formulas are given for calculating the number, average mol. weight and the average degree of polymerization from the nitrogen and hydroxyl group content. The optimal conditions of hardening of lacquer films and optimal NCO:OH ratios in the composition were investigated, and it was found that the rate of hardening decreases with increasing degree of polymerization. The relationship between the hardening of films and the average degree of polymerization and hardening temperature shows that by heating for 3 hours at 100C a 98 - 100% hardening is obtained for all lacquer coatings. At 90C the rate of hardening varies considerably in relation to the structure of the polymer, especially in the first hour of hardening. The coatings containing tetrafunctional polymers have a high gloss and hardness and excellent adhesion. The physico-mechanical properties and chemical stability are tabulated. All polyurethan coatings have good chemical stability. Orig. art. has: 5 figures and 4 tables.

ASSOCIATION: None

SUBMITTED: 00

ENCL: 00

SUB CODE: MT, OC

NO REF SOV: 005

APPROVED FOR RELEASE: 08/23/2000

OTHER: 002

Card 2/2

CIA-RDP86-00513R001652510009-2"

L 29997-65 EWT(m)/EPF(c)/EPR/EWP(j)/T Pc-L/Pr-L/Ps-L WW/RM ACCESSION NR: AP4047676 S/0303/64/000/005/0015/0019

21

AUTHOR: Sorokin, M.F.; Khinchina, E.L.

TITLE: The synthesis of linear polyesters from monoepoxide compounds and dicarboxylic acids. \(\lambda\)

SOURCE: Lakokrasochny\*ye materialy\* i ikh primeneniye, no. 5, 1964, 15-19

TOPIC TAGS: polyesterification, linear polyester, monoepoxide, dicarboxylic acid, molecular weight distribution, turbidimetric titration, adipic acid, glycidol ester, two stage synthesis, polydispersity

ABSTRACT: A new type of linear polyesters was synthesized from dicarboxylic acids and monoepoxide compounds without catalysts. The resulting resins were very soluble in aromatic hydrocarbons, ketones, acetates, chlorinated hydrocarbons and dioxane. The synthesized polyesters were analyzed for acid and hydroxyl numbers, reduced viscosity, molecular weight (by the methods of differential ebulliometry, cryoscopy, and terminal carboxyl and hydroxyl group determinations), and molecular weight distribution (by turbidimetric titration). The polyester synthesis process can be divided into two stages; the formation of acidic monoesters and the formation of diesters. A series of experiments was run with different initial concentrations of adipic acid and glycidol esters to determine

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L 29997-65

ACCESSION NR: AP4047676

the overall sequence of the synthesis stages. It was established that the speed constants are fixed and do not depend on the ratio of the original components, and that the speed of polyesterification increases with an increase of glycidol esters in the reaction mixture. The use of catalysts accelerates only the first stage of the synthesis. The properties of polyesters synthesized at a 1:1 ratio of the original components are presented. The effects of the types of epoxy compounds and dicarboxylic acids on the degree of conversion and molecular weight distribution of the polyester are shown. The molar ratio of the initial influence on the molecular weight distribution of the second stage of synthesis are analyzed for their influence on the molecular weight distribution of the polyesters. Orig. art. has: 5 figures, 4 tables, and 3 chemical equations.

ASSOCIATION: none

SUBMITTED: 00

ENCL: 00

SUB CODE: OC, MT

NO REF SOV: 002

OTHER: 005

Card 2/2

L 25065-65 EWT(m)/EPF(c)/EWP(j)/T Pc-4/Pr-4 RM

ACCESSION NR: AP5002211 5/0303/64/000/006/0001/0004

AUTHOR: Sorokin, M. F.; Khinchina, E. L.

TITLE: Coatings based on polyesters of momepoxy compounds and dicarboxylic acids

SOURCE: Lakokrasochnyye materialy i kh primeneniye, no. 6, 1964, 1-4

TOPIC TAGS: polyester coating, polymer coating, monoepoxy ester, dicarboxylic acid ester, coating hardness, coating elasticity, epoxy resin, allylglycidyl ether

ABSTRACT: A series of tests were carried out on the physical and chemical properties of coatings based on synthetic polyesters and E-40 epoxy resin; as well as on unsaturated linear polyesters from allylglycidyl ether with dicarboxylic acids. First, 30% solutions of polyesters and E-40 resin were mixed with equal parts of toluene, butyl acetate and butyl alcohol to form a mixture which can be kept for some time at room temperature without solidifying. When desired, it can be solidified in 3 hours at 150C. A full-page table lists 15 such compounds with their E-40 content and values for hardness, elasticity, and resistance to water, acids and alkalies. All coatings showed good luster and adhesion to metals, glass wood, but polyester coatings based on E-40 resin showed the highest elasticity /5

L 25065-65

ACCESSION NR: AP5002211

and impact toughness. Those with E-40 and phenylglycidol or mixed phenols had a hardness of 0.7 to 0.9, while those with allylglycidyl ether and dicarboxylic acids were very soft. Polyester E-40 coatings were then tested for aging at 60C, but showed no loss of elasticity or hardness after 500 hours. Coatings based on the unsaturated polyesters of allylglycidyl ether and dicarboxylic acids were also tested after drying for 2 hours with a desiccant of lead and manganese resinates. Double coatings 25-35,4 thick were tested after standing at room temperature for 7 days and showed an impact toughness of 50 kg/cm<sup>2</sup> and an elasticity of 1 mm. They showed no deterioration after submersion in 10% NaOH for 6 days. Orig. art. has: 2 tables and 3 figures.

ASSOCIATION: none

SUBMITTED: 00

ENCL: 00

SUB CODE: MT

NO REF SOV: 001

OTHER: 000

Card 2/2

SOROKIN, M.F.; LATOV, V.K.

Application of the light diffusion method for determining the molecular weights of polymers. Plast.massy no.6:49-51 '64. (MIRA 18:4)

ACCESSION NR: AP4037273

s/0190/64/006/005/0791/0797

AUTHORS: Kochnov, I. M.; Sorokin, M. F.

TITLE: Polymerization kinetics of glycidyl methacrylate

SOURCE: Vy\*sokomolekulyarny\*ye soyedineniya, v. 6, no. 5, 1964, 791-797

TOPIC TAGS: polymerization kinetics, methacrylate, azoisobutyric acid, initiator, polymer viscosity, molecular weight, rate constant, monomer

ABSTRACT: The kinetics of glycidyl methacrylate (GMA) radical polymerization was investigated in the presence of dinitryl azoisobutyric acid and benzoyl peroxide initiators. An empirical equation was established relating the polymer viscosity to its molecular weight, or  $\{\eta\} = 7.83 \cdot 10^{-3} \cdot M^{0.18}$ . From the experimental data the following values are obtained for initiator rate constants as functions of temperature  $k_p = 4.13 \cdot 10^5 \cdot \exp(-4000/RT)$ , and for the effective activation energy,

 $k_{\rm o}=6.17\cdot 10^7\cdot \exp\left(-400\,/\,RT\right)$  E = 18.8 kcal/mol. Other constants, such as the rate of growth termination and propagation of the chain through the monomer, have also been calculated. A list is made of GMA polymerization kinetics magnitudes with corresponding values for various

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ACCESSION NR: AP4037273

other ester methacryl acids. There is a close correspondence between the GMA results obtained by the authors and those methacryl esters with carbon atom numbers ranging from 1 to 4. Orig. art. has: 9 formulas, 4 figures, and 2 tables.

ASSOCIATION: Moskovskiy khimiko-tekhnologicheskiy institut im. D. I. Mendeleyeva (Moscow Institute of Chemical Engineering)

SUBMITTED: 28May63

DATE ACQ: 09Jun64

ENCL: 00

SUB CODE: OC

NO REF SOV: 005

OTHER: 016

Card 2/2

APPROVED FOR RELEASE: 08/23/2000 CIA-RDP86-

CIA-RDP86-00513R001652510009-2"

ACCESSION NR: AP4037274

5/0190/64/006/005/0798/0802

AUTHORS: Sorokin, M. F.; Kochnov, I. M.

TITLE: Relative monomer reactivity in copolymerization of glycidyl methacrylate with styrene

SOURCE: Vy\*sokomolekulyarny\*ye soyedineniya, v. 6, no. 5, 1964, 798-802

TOPIC TAGS: monomer reactivity, copolymerization, glycidyl methacrylate, styrene, initiator, benzoyl peroxide, reactivity ratio

ABSTRACT: Copolymerization of glycidyl methacrylate (GMA) with styrene in the presence of recrystallized dinitryl azoisobutyric acid and benzoyl peroxide has been carried out in various solvents at 60 and 120C in a nitrogen atmosphere. The composition of the resultant copolymers and the monomer reactivity ratios  $r_1$  and  $r_2$  have been calculated by the method of M. Fineman and S. D. Ross (J. Polymer Soi., 5, 259, 1950), producing  $r_1$  = 0.55 and  $r_2$  = 0.45. The monomer reactivity ratios are shown to be independent of the nature of solvent and initiator used. The variation of  $r_1$  and  $r_2$  with the temperature is given by an expression of the form

 $r = \exp(-\Delta E/RT)$ , tending to unity with increase in temperature. The parameters Q and e in the Card 1/2

SOROKIN, Mikhail Fedorovich; LYALYUSHKO, Kapitolina Alekseyevna; YUKHNOVSKIY, G.L., prof., doktor khim. nauk, retsenzent; ARKHIPOV, M.I., doktor tekhn. nauk, prof., retsenzent; ALAVERDOV, Ya.G., red.

[Practical laboratory work on synthetic polymers for lacquers] Praktikum po sinteticheskim polimeram dlia lakov. Moskva, Vysshaia shkola, 1965. 271 p. (MIRA 18:7)

1. Zaveduyushchiy kafedroy Khar'kovskogo Politekhnicheskogo instituta im. V.I.Lenina (for Yukhnovskiy).

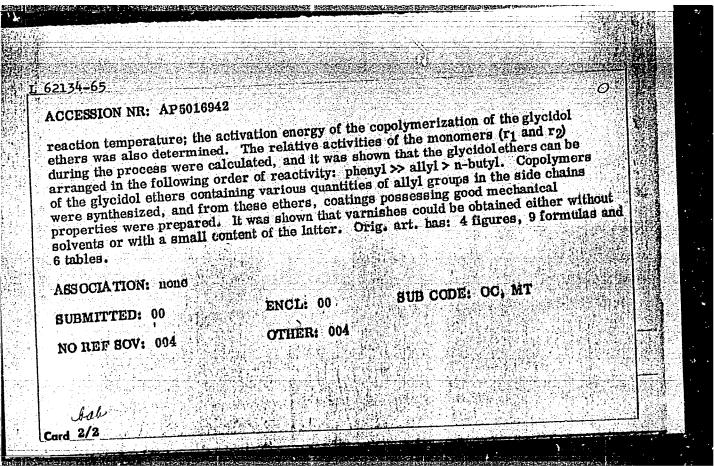
EWG(j)/EWT(m)/EPF(c)/EPF(n)-2/EWP(j)/T/EWA(h)/EWA(1) Pc-4/ L 54610-65 Pr-4/Peb/Pu-4 GG/RM ACCESSION NR: AP5011238 UR/0303/65/000/002/0001/0005 Sorokin, M. F.; Lyalyushko, K. A.; T'ang, Tsung-lan Effect of gamma radiation on certain epoxy resins Lakokrasochnyye materialy i ikh primeneniye, no. 2, 1965, 1-5 epoxy resin, gamma radiation, radiation resistance, TOPIC TAGS: coating ABSTRACT: A study has been made of 1) the effect of y-radiation on the properties of certain epoxy resins, and 2) the effect of certain hardeners on the radiation resistance of the epoxy coatings. The resins used were based on epichlorhydrin and. 2, 2-bis (4-hydroxyphenyl) propane or -butane, or 1,1-bis(4-hydroxyphenyl)cyclohexane. The resins were irradiated from a Go o source in glass test tubes and changes in solubility in organic solvents, melting or dropping point, glycidyl group content, molecular weight, intrinsic viscosity, poly-dispersity and weight loss at 250C were determined. The data are presented in graphic and tabular form. It was found that at doses up

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dose is raised to	properties remain virtual 775 Mrad, the glycidyl groweight and intrinsic viscolose. The mechanical properties	oup content drops linea osity rise logarithmics	11 <b>y</b>
of coatings of the	same resins were tested	in the presence of such nodiphenylmethane, hexa	<b>.</b>
and	nd a nolvamida ragin. V in	B CHES KIASH IN CORRE	🕨 e i grafija i grafija 🕳 e i grafija 🕳 e i grafija i
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form, indicate the suitable as radiat 6 tables and 2 for ASSOCIATION: none	ind a polyamide resin. Vine it the films show good rad: ion-resistant coatings; mulas.	iation resistance and a Orig. art. has: 7 figu [SM]	re res,

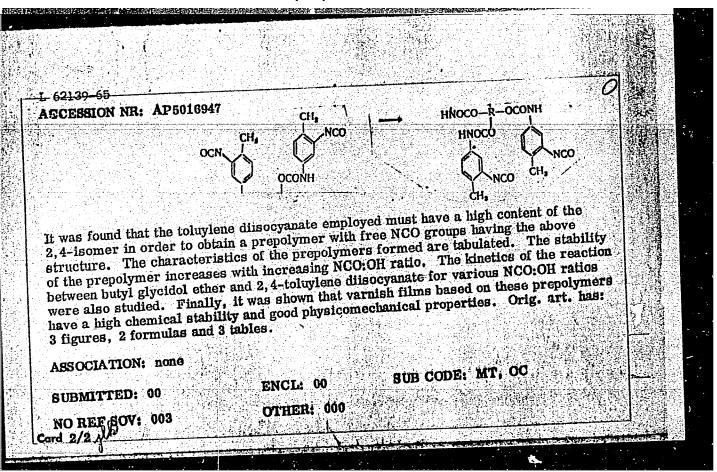
SOROKIN, M.F.; LATOV, V.K.

Ebulliometric determination of the molecular weight of polymers. Plast. massy no.2:54-58 '65. (MIRA 18:7)

62134-65 EWT(m)/EPF(c)/EWP(1)/ED	PR/EWP(j)/T/EWP(t)/EWP(b) -Pc-4/Pr-4/ UR/0303/65/000/003/C001/0007
4 RPL JD/WW/RM ACCESSION NR: AP5016942	667.633.263.3
AUTHOR: Sorokin, M.F.; Kochnov, I.M	.; Krivopalova, I.S.
TITLE: Synthesis of copolymers of aryl	and alkyl glycidol ethers and the preparation
at anatings from them	언니는 하루는 공장은 경험을 통한 사람이 하는 경험을 받았다. 이 사람이 아름다는 경험을 들어 보다면 모양이 가는 그는 그는 사람들이 가득하는 살아지를 받았다면 목표를 했다.
SOURCE: Lakokrasochnyye materialy i TOPIC TAGS: glycidol ether, protective	coating, copolymerization, polyether
ABSTRACT: The authors studied the co- ether with allyl glycidol ether in the pre- copolymers obtained and the coatings ba	polymerization of phenyl and nonperties of the sence of alkali catalysts, the properties of the sed on them. The kinetics of this copolymerizased on them. The kinetics of this copolymerization and it was shown that the process occurs via
a stepwise ionic mechanism at the ends of the polymer chains. The	mation of polyethers having hydroxyl global enters process of copolymerization of glycidol ethers of the nonstationary period (C) and overall rate of the nonstationary period and rate constant of chain growth stationary period and rate constant of chain growth of monomers in the reaction mixture and on the
were shown to depend of the condition of	



L 62139-65 EWT(m)/EWP(j)/T Pc-	-4 RM (200 (2002 (2003)	
ACCESSION NR: AP5016947	UR/0303/65/000/003/0032/0035 /6 667.633.263.3	
AUTHOR: Sorokin, M.F.; Kochnova,	Z.A.; Zhukova, L.B.	
TITLE: One-component polyurethand functional polymers of glycidol ethers	y mighed hased on ni rogen-containing terra-	
Torogram Takokragochnyve material	ly i ikh primeneniye, no. 3, 1965, 32-35	
TOPIC TAGS: polyurethane varnish,	, glycidol ether, toluyelene diisocyanate, nitrogenous	を受ける。
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nitrogen-containing polymore from the	e stable polyurethane varnishes based on tetrafunctional did not be stable polymers with side of the stable polymers with the see polymers and toluylene discoyanate. The reaction are polymers and toluylene discoyanate.	
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L 65218-65 EWT(m)/EPF(c)/EWP(j)/T RM

ACCESSION NR: AP5022506

UR/0303/65/000/004/0004/0007 666.21.633

AUTHOR: Sorokin, M. F., Lyalyushko, K. A.; Belokrinitskaya, N. Ye.

TITLE: Epoxy resins based on dicyclopentadiene and polyhydric alcohols

SOURCE: Lakokrasochnyye materialy i ikh primeneniye, no. 4, 1965, 4-7

TOPIC TAGS: epoxide, protective coating, vernish, polymerization

ABSTRACT: D Alicyclic diepoxides have been prepared which are suitable for making varnish films having good mechanical properties. It is noted that alicyclic epoxides are of interest because the raw materials for them are readily available, and because films from them show improved light and thermal stability and better electrical insulation properties. First, bis(dicyclopentadienyl) ethers of ethylene glycol, diethylene glycol, and glycerol were synthesised in the presence of H2SO, in 70—80% yields. From the ethers the diepoxides were obtained by epoxidation with peracetic acid in 70—80% yields. Both the ethers and the diepoxides were suitable for preparing varnish films with good mechanical properties. The ethers polymerized via the dicyclopentadienyl double bond in the presence of driers to form varnish films with an impact strength of 50 kg.cm. The diepoxides in the form of 60% solutions in

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ACCESSION NR: AP5022506			
cellusolve cured in the presentilms had an impact strength out degradation. Orig. art.	of 10-50 kg · cm and vitustood	COOC 110M TO	
ASSOCIATION: none			
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SOROKIN, M.F.; BABKINA, M.M.

Composition of tricopolymers and the copolymerization constants of butyl methacrylate, glycidyl methacrylate, and methacrylic acid. Vysokom. soed. 7 no.4:737-740 Ap '65. (MTRA 18:6)

1. Moskovskiy khimiko-tekhnologicheskiy institut imeni Mendeleyeva.

KOCHNOV, I.M.; SOROKIN, M.F. Kinetics of copolymerization of glycidyl methacrylate with styrene. Vysokom. soed. 7 no.11:1916-1922 N 165.

(MIRA 19:1)

1. Moskovskiy khimiko-tekhnologicheskiy institut imeni D.I. Mendeleyeva. Submitted December 11, 1964.

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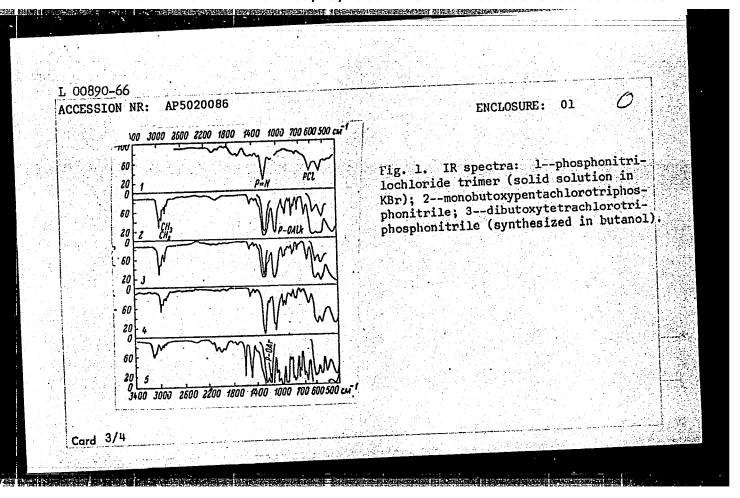
SOROKIN, M.F.; BAHKINA, M.M.

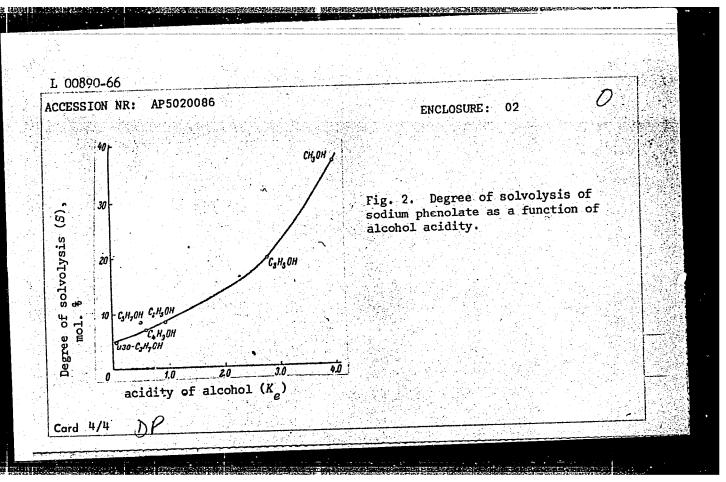
Fractionation of a triple copylymer of butyl, methacrylate, clycidyl methacrylate, clycidyl methacrylate, and methacrylic acid. Vysokom. soed. 8 no. 1:115-119 Ja \*66 (NTRA 19:1)

1. Moskovskiy khimiko-tekhnologicheskiy institut imeni Mendeleyeva. Submitted February 24, 1965.

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UTHOR: Sore	okin, M. F.; Lator	v, v. K. 44,55		1	39	
ITLE: Synt	nesis of partial o	esters of phosp	honitrilochlor	<u>ide</u> \trimer	B	
OURCE: Zhu	rnal obshchey khi	mii, v. 35, no.	8, 1965, 1471 55	_1475		
TOPIC TAGS:	ester, phosphoni	trile, polymer,	chlorinated a	liphatic compound	1	
	onobutoxypentach1	orotriphosphoni	trile, dibuto	ytetrachiorotrip	lospilo-	
itrile		C4H0O II L	,0C₄H₀			
		CI/P\N	\CI			
and tributox	ytrichlorotriphos	CI P OC	H <sub>0</sub>			
		C4HOO P	`0C¹II°			

L 00890-66 ACCESSION NR: AP5020086  were synthesized from phosphonitrilochloride trimer and sodi alcohol-benzene solvent. The respective yields were 58.3, 5 alcohol-benzene solvent, refractive indices, and chemical points, specific gravities, refractive indices, and chemical points, specific gravities, Diphenoxytetrachlorotriphospections.	t townii ac Mei.e nerer
mined for all these compounds. Supplied these four compaired, in the 60% yield, from phosphonitrilochloride and social alcohol-benzene solvent. The IR-spectra for these four compaired to spectrophotometer, are shown in fig. 1 of the Enclosure. 10 spectrophotometer, are shown in fig. 1 of the Enclosure. Sis (S) of sodium phenolate in reaction of the phosphonitric sodium phenolate as a function of the acidity of alcohol so sodium phenolate as a function of the acidity of alcohol so sodium phenolate. It was found that during solvolys fig. 2 of the Enclosure. It was found that during solvolys in alcohol, substitution with alkoxy-group takes place side in alcohol, substitution of the acidity of alcohol solve the phenoxy-group. Orig. art. has: 2 figures and 1	dium phenolate using an pounds, taken with a UR- The degree of solvoly- lochloride trimer with lvent (K) is shown in is of sodium phenolate by side with substitutable.
ASSOCIATION: Moskovskiy khimiko-tekhnologicheskiy institut (Moscow Institute of Chemical Technology)	: im. D. I. Mendeleyeva
	SUB CODE: GC, OC
SUBMITTED: 08Jun64 ENCL: 02	
SUBMITTED: 08Jun64 ENCL: 02  NO REF SOV: 003 OTHER: 014	





# "APPROVED FOR RELEASE: 08/23/2000 C

CIA-RDP86-00513R001652510009-2

UR/0303/65/000/005/0012/0014 WW/RM EWT(m)/EWP(j)/T L 18013-66 SOURCE CODE: AP6004313 ACC NR: Sorokin, M. F.; Babkina, M. M. AUTHOR: TITLE: Film-forming properties of ternary copolymers of butyl methacrylate, gly-ORG: none cidyl methacrylate, and methacrylic acid SOURCE: Lakokrasochnyye materialy i ikh primeneniye, no. 5, 1965, 12-14 TOPIC TAGS: methacrylate plastic, copolymer, thermosetting material ABSTRACT: The film-forming properties of ternary copolymers of butyl methacrylate (BMA), glycidyl methacrylate (GMA), and methacrylic acid (MAA) were investigated with the aim of developing new thermosetting methacrylic copolymers. 20% lacquers were prepared from these copolymers in a mixture of solvents of the following composition: toluene, 30%; cyclohexanone, 30%; butyl acetate, 14%; acetone, 26%. Epoxy resins E-40 and E-181 were used to modify the lacquer films. The films were deposited on metal surfaces. The measured film-forming properties are tabulated. The composition of the copolymer was found to affect the properties of the lacquer films: as the content of reactive groups increases in the copolymer, the flexibi-UDC: 667.633.263.3 Card 1/2

,	8013-66 NR: AP60	04313					
lity tran	declines	, and the	t is the lack	chemical stabil dare capable of of impact streng phthalate (DBP) ining the same.	th. When the	- impact strengt	th
tabl	cope: 07		SUBM DATE: 00			OTH REF: 000	

SOURCE CODE: UR/0190/66/008/001/0115/0119 AP6003422 ACC NR: AUTHORS: Sorokin, M. F.; Babkina, M. M. ORG: Moscow Institute of Chemical Engineering im. D. I. Mendeleyev (Moskovskiy khimiko-tekhnologicheskiy institut) TITLE: Fractionation of a triple copolymer of butyl methacrylate, glycidyl methacrylate, and methacrylic acid SOURCE: Vysokomolekulyarnyye soyedineniya, v. 8, no. 1, 1966, 115-119

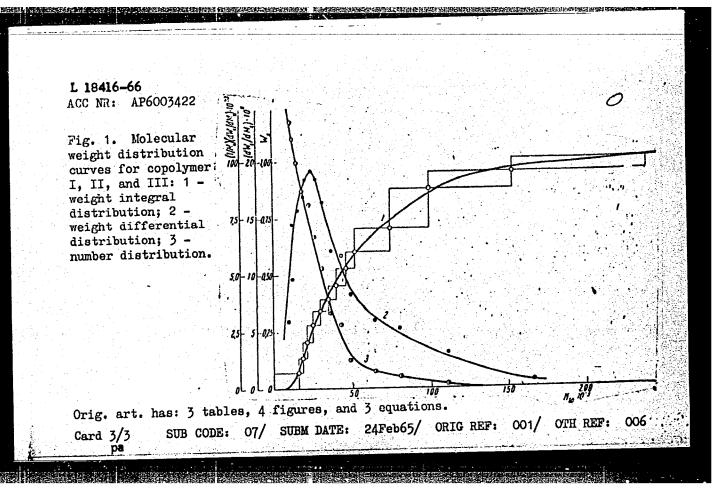
copolymer, methacrylate plastic, polymerization degree TOPIC TAGS:

ABSTRACT: Molecular weight distribution of a triple copolymer of butyl methacrylate (I), glycidyl methacrylate (II), and methacrylic acid (III) was investigated by means of fractionation and examination of properties of the obtained fractions. Composition of the copolymer in mole % is: I:II:III = 74.10: 9.65: 16.25. Synthesis followed that described previously by the authors (Tr. Mosk. khim.-tekhnol. in-ta im. D. I. Mendeleyeva, vyp. 48, 1965). Fractionation was achieved by means of a repeated fractional precipitation with water from 2% solution in dioxane at 20C. Molecular weight, composition, and specific viscosity of the

Card 1/3

UDC: 678.01:53+678.744

AC CC	polymer	were de	termined i	n each fra	ction. Mo	lecular we of concent stant K' =	ight distri ration was . 0.95 was c	bution curve determined alculated wh acetone solven by the ec	s ich
a. p	ermitted	to the determine empiri	nation of cal relati	n at cert on between	ain copoly η and mo	mer concer olecular we	ntrations in eight is giv	acetone solven by the ec	lu-

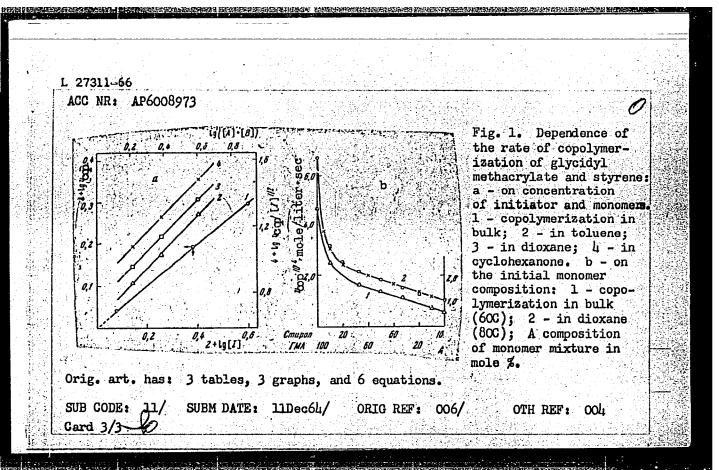


EWT(m)/EWP(i)/T/ETC(m)-6 LJP(c) WW/RM SOURCE CODE: UR/0190/66/008/003/0444/0449 AP6010108 ACC NR Sorokin, M. F.; Manovichu, I. AUTHORS: ORG: Moscow Chemical and Technological Institute im. D. (Moskovskiy khimiko-tekhnologicheskiy institut) of allyphosphinic and dichloride with glycols TITLE: Polycondensation and distomic phenols SOURCE: Vysokomolekulyarnyye soyedineniya, v. 8, no. 3, 1966, 444-449 TOPIC TAGS: glycol, phosphorus chloride, diethylene glycol, hydroquinone, resorcinol, hydrogen chloride, propane, phenol, polycondensation ABSTRACT: Self-extinguishing phosphorus containing polyesters have been prepared by polycondensation of allyphosphinic acid dichlorides with diethylene glycol, triethylene glycol, hydroquinone, resorcinol, and 2.2-di-(4-hydroxyphenyl) propane. It was established that oligomers are formed during the interaction of acid dichlorides of allylphosphinic acids with the dihydroxy compounds. Parallel to the formation of polyesters from glycols, the polymer decomposition caused by the hydrogen chloride liberated during the reaction, occurs; there is no decomposition with the use of diatomic phenols. It was found that phosphorus-containing polyesters are not easily inflammable and UDC: 541.64+678.86 Card 1/2

extinguish instar	atly after being taken out	of the flame. Orig.	art.
has: 3 figures ar	nd 2 tables. [Based on aut]	nor's abstract]	[NT]
SUB CODE: 07/	SUBM DATE: 25Mar65/ OTH REF: 014/	orig ref: 007/	
	도 한 시간 중에 당한 분들이 하고 있는 것을 하는 것이다. 이 1일 1명 회사를 하고 하는 것 같다고 하고 있다.		
	- 그는 이 교육 관련적 취임 등에 다른 보다 되었다. 		
		- 이 제 로그 보고 1개를 통해될 수 있다. 	
<u>p</u>			
Card 2/2 UV			

L 27311-66 : EWT(m)/EWP(j)/T   IJP(c)   WW/RM	
ACC NR: AP6008973 SOURCE CODE	: UR/0190/65/007/011/1916/1922
AUTHORS: Kochnov, I. M.; Sorokin, M. F.	26 B
ORG: Moscow Institute of Chemical Technology im. I khimiko-tekhnologicheskiy institut)	D. I. Mendeleyev (Moskovskiy
TITLE: Kinetics of copolymerization of glycidyl me	ethacrylate and styrene
SOURCE: Vysokomolekulyarnyye soyedineniya, v. 7, r	10. 11, 1965, 1916-1922
TOPIC TAGS: copolymer, radical polymerization, pol	Lymerization kinetics, styrene
ABSTRACT: The effect of different solvents and inj	tiators on the radical
copolymerization kinetics of the reaction between g styrene was determined. The reaction was studied i	glycidyl methacrylate and
toluene, cyclonexanone, and dioxane. Diniryl of ag	chutyric acid and hengori
peroxide were used as initiators. The copolymeriza	tion rate constants the rote
of copolymerization, and the cross termination consfunctions of the initial concentration of reactants	tants p were determined as
initiator. The activation energy of the conclumeri	zation was also determined.
ne rate of the reaction obeyed the expression	
Gard 1/3	DC: 66.095.26+678.744+678.746

L 27311-66	
ACC NR: AP6008973 $v_{\text{COD}} = K_{\text{COD}}[1]^{\frac{1}{1}}([A] + [B])_{11}$	7
where v <sub>cop</sub> is the rate of copolymerization, K <sub>cop</sub> is the rate constant (I), and	
((A) + (B)) the initiator and total monomer concentration respectively. The crostermination constants were derived after I. M. Kochonov (Dissertatsiya, 1964)	8
$\int_{\rho} \frac{\overline{\nu_{\mathbf{A}} \cdot \nu_{\mathbf{B}} \cdot \psi}}{(\rho + 2 \cdot \Phi \cdot \gamma + \mu)^{\frac{1}{N}}} \cdot \sqrt{\frac{\rho}{2}}$	
where $\psi = r_1[A]^2 + 2[A] \cdot [B] + r_2[B]^2$ , $\rho = (r_1 \cdot \nu_B \cdot [A_p] \cdot [A])^2$ , $\gamma = \frac{r_1 \cdot r_2 \cdot \nu_B[A_p][B_p][A][B]}{\equiv r_1 \cdot r_2 \cdot \nu_B[A_p][B_p][A][B]}$ and $\mu = (r_2 \cdot \nu_A[B_p][B])^2$ ; $[A_p]$ and $[B_p]$	
where $v_A$ and $v_B$ are the separate polymerization rates for the polymerization of $\Lambda$	
and B respectively, and (Ap), (Bp), (A), (B) are the initial monomer concentrations for the individual and copolymerization of A and B respectively. The experimental results are presented in graphs and tables (see Fig. 1). It was	
found that the energy of activation for the copolymerization was independent of the nature of the solvent.	
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는 이 사용 전 경험 한 경험 수 있다. 이 사이 사용 전 경험 전 경험 등 경험 전 경험 등 하는 것이 되었다. 그런 것이 되었다. 그런 것이 되었다. 그런 것이 되었다. 그런 것이 없는 것이 되었다. 	
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UR/030 1/66/000/003/0003/0006 SOURCE CODE: ACC NR. AF6019445 (A) AUTHOR: Sorckin, M. F.; Sokol, I. N. ORG: none with esterified phenolformaldehyde Setting of epoxy resins TITLE: resins SOURCE: Lakokrasochnyye materialy 1 ikh primeneniye, no. 3, 1966, 3-6 TOPIC TAGS: phenolformaldehyde, phenolformaldehyde resin, epoxy Phenolformaldehyde resins esterified with allyl bromide can be used as hardening agents for epoxy resins, including low-molecular, They are found to be more active hardeners than nonesterified. phenolformaldehyde resins. Compound compositions with hardening properties can be produced without solvents. Films made of these compositions are found to have high mechanical properties. Orig. art has: 4 figures and 2 tables. [Translation of authors' abstract] SUBM DATE: none/ ORIG REF: 003/ SUB CODE: 07/ Card 1/1

L 29545-66 EWT(m)/EWP(j)/T IJP(c) RM

ACC NR: AP6007772 (N)

SOURCE CODE: UR/0195/66/007/001/0042/0048

AUTHOR: Sorokin, M. F.; Latov, V. K.

ORG: Moscow Chemical Engineering Institute im. D. I. Mendeleyev (Moskovskiy

khimikotekhnologicheskiy institut)

TITLE: Kinetics of the reaction of phosphonitrile chloride trimer with Na alcoholates

SOURCE: Kinetika i kataliz, v. 7, no. 1, 1966, 42-48

TOPIC TAGS: phosphonitrile, ethanol, methanol, chloride, sodium compound, reaction rate

ABSTRACT: The kinetics of the reaction of phosphonitrile chloride (PNC) trimer and its incomplete butyl esters (monobutoxypentachlorotriphosphonitrile, dibutoxytetra-chlorotriphosphonitrile, and tributoxytrichlorotriphosphonitrile) with sodium butoxide in butanol, sodium ethoxide in ethanol, and sodium methoxide in methanol were studied. The reaction is first order with respect to the PNC trimer, its esters, and the alkoxide ion, and zero order with respect to the alcohol. An

Card 1/2

UDC: 541.124 : 542.951.3

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L 29545-66

ACC NR: AP6007772

increase in the dielectric constant of the medium was found to accelerate the reaction rate. An increase in the degree of substitution of chlorine atoms in the PNC trimer by butoxy groups sharply lowers the reaction rate, raises the activation energy, and leads to a decrease in the entropy of activation and the steric factor. A transitional state involving a change from the tetra- to the pentacoordination of phosphorus is proposed. Orig. art. has: 5 figures, 3 tables, and 6 formulas.

SUB CODE: 07/ SUBM DATE: 12May64/ ORIG REF: 004/ OTH REF: 014

Card 2/2 W

ACC NRI AP6006717 (A) SOURCE CODE: UR/0303

SOURCE CODE: UR/0303/66/000/001/0004/0009

AUTHOR: Sorokin, M. F.; Chibisova, Ye. I.

ORG: none

TITLE: Polyester varmish resins containing anhydride of 1,4,5,6,7,7-hexachlorobicycle-(2,2,1)-5-heptyl-2,3-dicarboxylic acid

SOURCE: Lakokrasochnyye materialy i ikh primeneniye, no. 1, 1966, 4-9

TOPIC TAGS: phthalic anhydride, polyester plastic, varnish, paint, lacquer, polycon-densation, aliphatic alcohol, polyhydroxy aliphatic alcohol

ABSTRACT: Kinetics of the polycondensation of the anhydride of 1,4,5,6,7,7-hexachloro-bicyclo-(2,2,1)-5-heptyl-2,3-dicarboxylic acid with ethyleneglycol, glycerine, and xy-litol was studied in the 140-230°C range. The molar ratios of anhydride to glycol were of 1:1 to 1:2.5 and the reaction duration was 2-6 hr. The properties of the reaction products as coatings were compared to those based on phthalic anhydride. The rates of the polycondensation reaction and the energies of activation are graphed and tabulated. It was found that this polycondensation is a two-stage reaction. The first of these stages is fast, almost independent of temperature, and leads to the formation of the acidic esters. The second stage, highly temperature dependent, was found to be a second order reaction. The activation energies of the polycondensation reactions involv-

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ACC NR: AR6031252 (A) SOURCE CODE: UR/0081/66/000/011/S020/S020

AUTHOR: Sorokin, M. F.; Babkina, M. M.

TITLE: Synthesis and study of butylmethacrylate copolymers with glycidylmethacrylate and methacrylic acid

SOURCE: Ref. zh. Khimiya, Part II, Abs. 11S121

REF SOURCE: Tr. Mosk. khim. -tekhnol., in-ta im. D. I. Mendeleyeva, vyp. 48, 1965, 201-207

TOPIC TAGS: copolymer, methacrylic acid, butylmethacrylate, glycidylmethacrylate, thermosetting copolymers

ABSTRACT: The synthesis of butylmethacrylate copolymers (I) with glycidylmethacrylate (II) and methacrylic acid (III) was carried out in cyclohexanone (IV) and dioxane (V) at 70, 80, and 90C. Benzoyl peroxide (IV) and dinitrile of azodi-isobutyric acid (VII) in amounts of 0.1, 0.2, 0.4 and 0.8 mol % were used as initiators. The monomer concentration in the reaction mixture amounted to 20, 30, and 40% and the molecular ratios varied over a wide range. The copolymerization rate (CR) increased with an increase in (II) concentration, while (III) in concentration of less

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#### ACC NR: AR6031252

than 25 mol % was found to retard CR. However, an acceleration of CR was observed with an increase in the mole fraction of III, accompanied by an increase in viscosity of the solution, which gelatinizes on reaching a 70% conversion of the monomer. Such a phenomenon is explained by the capacity of III to form an H-bond with compounds containing carbonylic oxygen, which is also true for II, IV and III. An increase in reaction temperature and in VI and VII concentrations causes an increase of the rate of copolymerization and a decrease in the molecular weight of the copolymers. The CR, the molecular weight, and the yield of copolymers increase with an increase in concentration of the monomers in the solution. IV and V do not substantially affect CR. The polydispersion of copolymers increases with an increase of I content and also with increases in the reaction temperature and an increase of I content and also with increases in the reaction temperature and concentrations of VI and VII. VI and V do not affect the polydispersion. The triple thermosetting copolymers obtained are capable of self setting at higher temperatures. Some properties of the copolymers were determined. V. Agasantoyan. [Translation of abstract]

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SUB CODE: 07/

Cord 2/2

SOROKIN, M. G. --

"Method of Calculation and Investigation of the Operation of a Wind Motor With Large Initial Starting Moment." Cand Tech Sci, All-Union Sci-Res Inst of Electrification of Agriculture, Moscow, 1954. (RZhMekh, Oct 54)

Survey of Scientific and Technical Dissertations Defended at USSR Higher Educational Institutions (10)

SO: Sum. No. 481, 5 May 55

Results of preliminary tests of the 1-D-18 wind power plant. Izv.
AN Kazakh.SSR.Ser.energ. no.4/5:128-141 '54. (MLRA 9:5)

(Wind power)

scacifan, ". o.

"Biological and Morphological Variations in the Baccon Dog (Nyctereutes procyonoides grai), Which Is Being Acclimated in Kalininskaya Oblast." Cand Biol Sci, Kalinin State Pedagogical Inst, Kalinin, 1954. (RZhBiol, No 1, Jan 55)

Survey of Scientific and Technical Dissertations Defended at USSR Higher Educational Institutions (13) 80: Sum. 509, 29 Jul 55

SHAPOSHNIKOV, L.V., doktor biolog.nauk, prof.; GOLOVIN, O.V.,kand.biolog.nauk; SOROKIN, M.G., kand.biolog.nauk; TARAKANOV, A.D., starshiy prepodavatel. Prinimali uchastiye: V'IUNOV, V.N.; SOKOLOV, P.P., inzh.-ryboved; VIKTOROV, G.S., tekhn.red.

[Animal world of Kalinin Province] Zhivotnyi mir Kalininskoi oblasti. Kalinin, Kalininskoe knizhnoe izd-vo. 1959. 459 p.
(MIRA 13:10)

1. Nachalinik Kalininskogo oblastnogo upravleniya okhotnichiyego khozyaystva (for Viyunov). (Kalinin Province---Vertebrates)

Morphology and biology of the northern birch mouse (Sicista betulina Pall.) in Kalinin Province. Nauch. trudy Kal. otd. MOIP no.2:31-40 '60. (KALININ PROVINCE\_BIRCH MOUSE)

以2015年1月1日中央 1月1日 | 1月1日 |

Sinel'nikova, L.A. and Sorokin, M.I. AUTHORS: 68-58-3-16/22 TITIE:

Automatic Weighing of Coal Blend Charged into Ovens (Avtomaticheskoye vzveshivaniye ugol'noy shikhty)

PERIODICAL: Koks i Khimiya, 1958, Nr 3, pp 55 - 56 (USSR).

ABSTRACT:

ACT: An installation for the automatic weighing of the coal blend charged to ovens developed by KIP on the Kuznetsk Metallurgical Combine is described. There is 1 figure.

ASSOCIATION: Kuznetskiy metallurgicheskiy kombinat

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(Kuznetsk Metallurgical Combine)

Card 1/1

KARPOV, A.G.; SOROKIN, M.I.

Computers serve the economy. Stal' 23 no.12:1122-1123 D '63.
(MIRA 17:2)

1. Volgogradskiy metallurgicheskiy zavod "Krasnyy Oktyabr'".

SOROKIN, M. I. Cand Agr Sci -- (diss) "On the problems of the agricultural engineering and personal seed growing in bottom lands." Mos, 1957. 21 pp (Mos Order of Lenin Agr Acad im K. A. Timiryazev), 110 copies (KL, 3-58, 98)

-42-

USSR/Cultivated Plants - Potatoes. Vegetables. Melons.

М

Abs Jour

: Ref Zhur Biol., No 12, 1958, 53612

Author

: Sorokin, M.I.

Inst

Moscow Agricultural Academy imeni K.A. Timiryazev

Title

: Some Methods of Increasing the Potato Yield and of

Improving Its Quality on Flood Lands.

Orig Pub

Dokl. Mosk. s.-kh. akad. imeni. K.A. Timiryazeva, 1957

vyp. 28, 222-227

Abstract

This is a study of the effect of mellowing and moling the tillable horizon of bottom land soil to the depth of 25-27 cm, - on the yield and the planting qualities of potato tubers of the Lorkh and Epron varieties. The soil was plowed to the depth of 25-27 cm. With plowing and pulverizing, the increase in the yield of the Lorkh variety comprised 26 centners/ha or 16.5%. With plowing

Card 1/2

- 38 -

POLTAVSKAYA, Lyudmila Semenovna, kand.biolog.nauk; SOROKIN. Mikhail Iyanovich, kand.sel'skokhoz.nauk; KELIN, M., red.; CHIZHIKOVA, V., tekhn.red.

[Corn pests and diseases and their control in Mordovia] Vrediteli i bolezni kukuruzy i bor'ba s nimi v usloviiakh Mordovii. Saransk, Mordovskoe knizhnoe izd-vo, 1960. 47 p.

(MIRA 14:3)

(Mordovia--Corn (Maize) -- Diseases and pests)

SOROKIN, M.I., kand.sel'skokhozyaystvennykh nauk; KOVAL'CHUK, P.A., agronom

Effectiveness of supplementary spring fertilizing of perennial grasses. Uch. zap. Mord. gos. un. no.13:106-109 '60.

(MIRA 15:11)

l. Kafedra agronomii i pochvovedeniya Mordovskogo gosudarstvennogo universiteta.

(Mordovia--Grasses--Fertilizers and manures)

SOROKIN, M.I., kand.sel¹skokhozyzystvennykh nauk; SHURKIN, V.P. Pea cultivation in the Mordovian A.S.S.R. Zemledelie 24 no.4:

(MIRA 15:4) 36-39 Ap 162.

1. Mordovskaya respublikanskaya gosudarstvennaya sel!skokhozyzystvennaya stantsiya. (Mordovia-Peas)

CIA-RDP86-00513R001652510009-2" APPROVED FOR RELEASE: 08/23/2000

SOROKIN, M.M.

Propagandists of new ideas. Nauka i zhyttia 11 no. 4:28 Ap 161. (MIRA 14:5)

1. Predsedatel' gruppy Obshchestva dlya rasprostraneniya politicheskikh i nauchnykh znaniy USSR na Khar'kovskom turbinnom zavode im. Kirova. (Kharkov-Machinery industry-Technological innovations)

137-58-4-6402

Translation from: Referativnyy zhurnal, Metallurgiya, 1958, Nr 4, p 10 (USSR)

AUTHOR: Sorokin, M.M.

Improving the Quality of Zinc and Lead Concentrates by Separa-TITLE:

tion of Middlings (Uluchsheniye kachestva tsinkovykh i svintso-

vykh kontsentratov s vydeleniyem promproduktov)

PERIODICAL: Sb. tr. Vses. n.-i. in-ta tsvetn. met., 1956, Nr 1, pp 17-29

Experiments performed on the removal of lead from Zn con-ABSTRACT:

centrate in the course of an industrial process have shown that removal of a portion of the Pb and Cu from the concentrates may be accomplished by refletation. The conditions of refletation depend upon the nature of the relationship among the Pb, Cu and Zn minerals. If these minerals are segregated, for the most part, separation of the Pb minerals may be performed after treatment of the concentrate by elevated amounts of a mixture of

cyanide and chlorine salts of calcium and sodium. If the Pb minerals are primarily in concretions with ZnS, separation requires very fine grinding of the concentrate. The best resu

in removal of zinc from Pb concentrate are obtained by de;

the PbS by lime (4.5-6 kg/t). The employment of CuSO<sub>4</sub> inc \_\_\_\_es Card 1/2

137-58-4-6402

Improving the Quality of Zinc and Lead Concentrates (cont.)

extraction of zinc in the froth product. As CuSO<sub>4</sub> consumption is varied from 0 to 2.0 kg/t, the extraction of Zn in the crude froth product rises from 30 to 75 percent, and the recovery of Pb from 10 to 20 percent. Investigation of the conditions for fining flotations of the froth product showed the best results to be obtained by fining the crude froth product with potassium bichromate.

A. Sh.

1. Ores--Processes 2. Ores--Separation 3. Flotation--Applications

Card 2/2

#### CIA-RDP86-00513R001652510009-2 "APPROVED FOR RELEASE: 08/23/2000

SOROKIN, M.M. pa

81998

s/120/60/000/03/038/055 E032/E514

AUTHOR:

The NVO-40 Air-Cooled High-Vacuum Oil Diffusion Pump

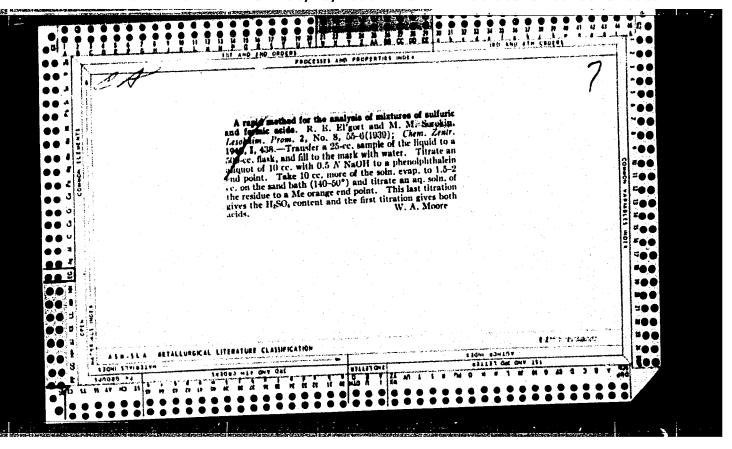
PERIODICAL: Pribory i tekhnika eksperimenta, 1960, No 3,

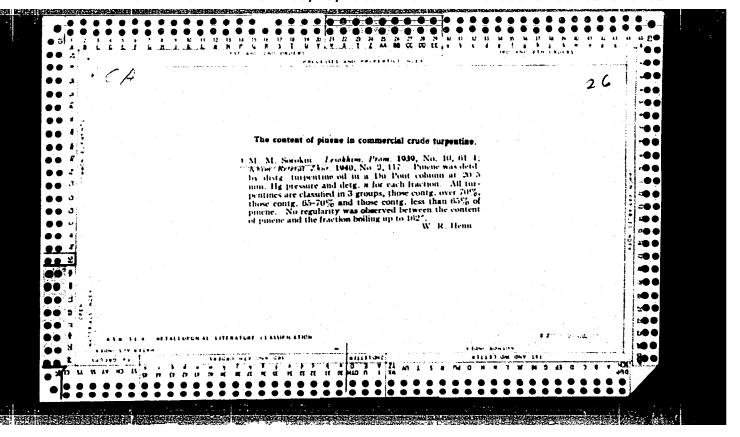
pp 131-133

ABSTRACT: A three-stage oil diffusion pump of an almost conventional design is described. A sectional drawing through the pump is shown in Fig 1. cooled with the aid of a built-in fan. The cooling is facilitated by a large number of cooling fins mounted on the outside of the pump. The design of the vapour column is shown in Fig 2. The overall dimensions of the pump are: length 28.2 cm, diameter 12 cm. characteristics are as follows: limiting vacuum  $5 \times 10^{-6}$  mm Hg, power consumption 0.45 kW, oil charge 100 cm3, mean pumping speed for air 50 l/sec, mean pumping speed for helium 100 &/sec (stable). weight of the pump is 8 kg. Fig 3 shows the pumping speed for air (1) and helium (2) as a function of the power consumption. Fig 4 shows the dependence of the pumping speed for air (1) and

Card 1/2

SORO	OKIN, M.M.		<b>V</b> • ***		
		promote advanced experience. Mashinostr (Kharkov-Turbogenerators)	roitel <sup>‡</sup> no (Mi	.7:24 J1 RA 15:7)	





SOROKIN, M.M. (Moskva)

Electrochemical disintegration of copper sulfides. Izv.AN SSSR.
Otd.tekh.nauk.Met.1 topl. no.3:104-108 My-Je '50.
(MIRA 13:6)

(Copper sulfides-Electrometallurgy)

GLEMBOTSKIY, V.A., doktor tekhn.nauk; SOROKIN, M.M., aspirant

Deactivation of sphalerite in an acid medium. Nauch.soob.
Inst.gor.dela 6:77-84 '60.

(Sphalerite)

GLEMBOTSKIY, V.A.; SOROKIN, M.M.

New inhibitor for bornite and chalcosine in the selection of combined copper-zinc and copper-lead concentrates. Dokl. AN SSSR 134 no.5: 1146-1149 0 160. (MIRA 13:10)

1. Institut gornogo dela Akademii nauk SSSR. Predstavleno akademikom A.A.Skochinskim.

(Bornite) (Chalcocite)

SOROKIN, M. M., Cand. Tech. Sci. (diss) "Investigation of Methods of Selection of Sphalerite and Secondary Sulfides of Copper under Conditions of Flotation of Copper-Zinc Ores," Moscow, 1961, 23 pp. (KrasnoyarskInst. Non-Ferr. Metals) 200 copies (KL Supp 12-61, 274).

SOROKIN, M.M., kand.tekhn.nauk, prof.; GLEMBOTSKIY, V.A., doktor tekhn.nauk; RAUKHVARER, Ye.L., kand.tekhn.nauk

Flotation properties of some compounds of the aromatic series. Nauch. soob. IGD 19:12-23 '63. (MIRA 17:2)

SOROKIN, M.M.; RAUKHVARGER, Ye.L.; SHCHEVELEVA, A.S.

Problem of the flotation action of willow oil and its components. Zhur. prikl. khim. 37 no.2:422-429 F '64.

(MIRA 17:9)

1. Institut gornogo dela imeni Skochinskogo.

(MIRA 18:1)

Decreasing the harmful effect of mineral salt in the flotation of martite. Gor.zhur. no.12:51-52 D 164.

1. Institut gornogo dela im. A.A.Skochinskogo.

10048/66/030/001/0037/0042 56	
SOURCE CODE: UR/004	
AUTHOR: Sorokin, M. M.; Baryshova, N. M.  AUTHOR: Sorokin, M. M.; Baryshova, N. M.  AUTHOR: obtaining ultrahigh vacuum with oil diffusion pumps (Transactions of the Physics of Thin Ferromagnetic vilus heid at Irkutsk)  Second All-Union Symposium on the Physics of Thin Ferromagnetic vilus heid at Irkutsk  TOFIC TAGS: ultrahigh vacuum, high vacuum technique, vacuum pump, diffusion pumps (vacuum oil  ABSTRACT: Examples of Soviet pumps employing type VM-1 oil diffusion pumps as to reach vacuum high vacuum the higher than 5x 10- tor at 20C) were used to illustrate to reach vacuum high vacuum high vacuum pumps to reach vacuum high vacuum hi	
and louvered after 24 no composition the press and investigated after and the composition the press and open manometer and the composition the press and the composition open manometer and the composition the press and the composition open manometer and the composition of pumping open manometer and the composition open manometer a	

ACC NR: AP600LL63:  $2 \times 10^{-8}$  torr. In the earlier higher vacuum period no molecules with molecular weight exceeding 44 were observed in the residual gas; after 30 hours heavier components appeared and ultimately predominated. In the absence of lower vapor pressure oils, such as the O8-124 and DC 705 fluids available in Western Europe and America, higher vacua can be reached only with the aid of liquid nitrogen cooled traps and sorption traps d Several such traps of Soviet manufacture are briefly described. With a type N-015-S pump and a liquid nitrogen or an activated charcoal trap, a vacuum of  $2 \times 10^{-8}$  torr was reached in which the residual gas was mostly hydrogen, contained nothing of molecular weight greater than 44, and remained unchanged in composition during 100 hours of operation. The types VA-05-5 and VA-5-7 vacuum systems (not described) performed considerably better, it being possible to achieve an oil-free vacuum of 10<sup>-10</sup> torr. It is cor luded that oil diffusion systems deserve more wide-spread use for the production of oil-free ultrahigh vacuum than they presently enjoy in the Soviet Union. Orig. art. has: 6 figures. SUB CODE: SUBM DATE: none/ ATD PRESS: 4206 Card 2/275

EWT(1)/EWT(m)/EPF(n)-2/T/ETC(m)-6L 16986-66 ACC NR: AP6001581 SOURCE CODE: UR/0120/65/000/006/0145/0149 Sorokin, M. M. AUTHOR: ORG: none TITLE: High-vacuum oil vapor pumps with improved characteristics SOURCE: Pribory i tekhnika eksperimenta, no. 6, 1965, 145-149 TOPIC TAGS: Vacuum pump, high vacuum pump, high vacuum technique / TaVL-1005 vacuum pump, N-1S-2 yacuum pump, N-5S-22 vacuum pump, N-2T-2 vacuum pump, N-5T-2 vacuum pump, N-6T-2 vacuum pump ABSTRACT: High-vacuum oil vapor pumps with improved characteristics are discussed. Most commercial vacuum pumps (such as TsVL-100S, N-2T, N-5T, N-8T) can operate at  $10^{-5}$  -- 2.10<sup>-4</sup> tor with a maximum exhaust pressure of 7.5·10<sup>-2</sup> -- 10<sup>-1</sup> tor. Some operating characteristics of a common pump (NVO-40) are presented as an example. By improving the cleaning and fractionation of the oil, by using metal seals, and by heating the chamber to 300-350C, the pump range can be improved to 3-5.10-8 tor. Based on the configuration shown in Fig. 1, a number of pumps have been Card 1/2 UDC: 533.563.5

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ACC NR: AP6001581

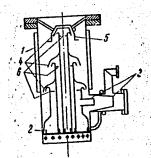


Fig. 1. Schematic diagram: 1 - shortened cooling jacket; 2 - fractionation rings; 3 - ejector; 4 - "umbrella" stages; 5 - oil baffle; 6 - throttling plates.

developed recently. These "Series 2" pumps with capacities of 100-8000 liter/sec have more desirable characteristics. All these pumps (N-1S-2, N-5S-2, N-5T-2, N-8T-2) are similar in construction to N-2T-2. A smaller capacity pump (15 liter/sec, 10-6 tor) was also developed. A table of the operating ranges of these vacuum pumps is presented. The author thanks K. A. Nesterovaya, N. Z. Sabirzyanov, F. D. Putilovskiy, I. M. Chistopolov, N. M. Mosin, and R. S. Gariffulin for their participation in the work. Orig. art. has: 6 figures and 1 table.

SUB CODE: 13/ SUBM DATE: 14Dec64/ ORIG REF: 002/ OTH REF: 002

Card 2/2 7285

ACC NRI AP6036839

SOURCE CODE: UR/0020/66/171/002/0320/0323

AUTHOR: Bokshitskiy, I. Ya.; Yelyutin, O. P.; Rogova, I. V.; Sorokin, M. N.

ORG: Central Scientific Research Institute of Ferrous Metallurgy im. I. P. Bardin (Tsentral'nyy nauchno-issledovatel'skiy institut chernoy metallurgii)

TITLE: Influence of group-IV transition elements and of Cu on the structure and physical properties of alloys based on the compound NiMn

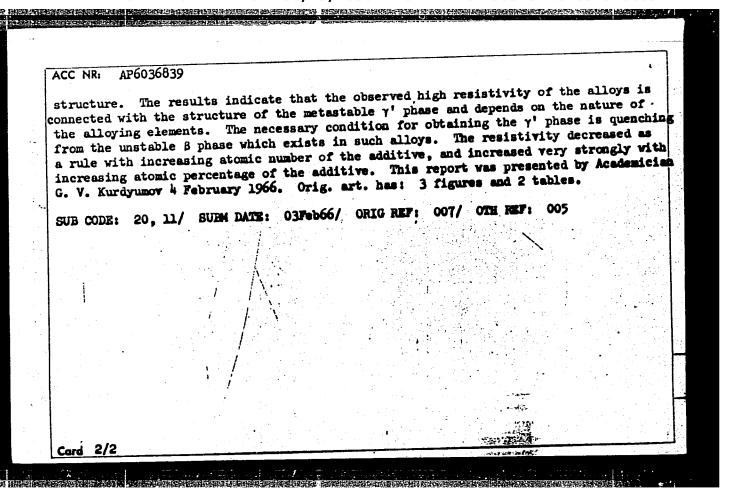
SOURCE: AN SSSR. Doklady, v. 171, no. 2, 1966, 320-323

TOPIC TAGS: nickel alloy, manganese containing alloy, transition element, alloy, phase diagram, resistivity, electric property

ABSTRACT: To investigate the structure and physical properties of alloys of the compound NiMn with transition elements, the authors fused pseudobinary alloys NiMn-Me (Ti, V, Cr, Fe, Co, Cu) containing 1-20 at.% V and Ti, and 1-10 at.% Cr, Fe, Co, Cu. The tests considered of a dilatometric analysis in the 100 -- 950° interval, measurements of the electric resistivity as a function of the alloying-additive content, an electron-microscopic investigation of the structure, and an x-ray phase analysis. The dependence of the electric properties and of the structure of the alloy as a function of the heat treatment was tested in the case of NiMn + 10 at.% V. The tests yielded the phase compositions of the different alloys and the types of crystal

Card 1/2

UDC: [669.018.5:669.017.11:537.3:669.017.3:621.78



KUNAKOV, Ya.N.; LIVSHITS, B.G.; SOROKIN, M.N.

Deformation of textures in silicon iron. Izv. vys. ucheb. zav.; chern. met. 6 no.5:146-150 '63. (MIRA 16:7)

1. Moskovskiy institut stali i splavov. (Iron-silicon alloys—Metallography) (Deformation (Mechanics))

L 13381-66 EWP(e)/EWT(m)/EWA(d)/EWP(t)/EWP(z)/EWP(b) IJP(c) JI ACC NR: AP6002905 SOURCE CODE: UR/0286/65/000/026	
ACC NR: AP6002905 SOURCE CODE: UR/0286/65/000/024	/0072/0072
INVENTOR: Yelyutin, O. P.; Bokshitskiy, I. Ya.; Rogova, I. Sorokin, M. N.	v.: 48
ORG: none	$\mathcal{B}$
TITLE: High-resistivity alloy, Class 40, No. 177075 [annual contents of the co	ounced by
Central Scientific Research Institute of Ferrous Metallurgy	im.
I. P. Bardin (Tsentral'nyy nauchno-issledovatel'skiy institute metallurgii)	ut Chernoy
SOURCE: Byulleten' izobreteniy i tovarnykh znakov, no. 24,	1965, 72
TOPIC TAGS: alloy, nigh resistivity alloy, nickel containing manganese containing alloy, vanadiar containing alloy	ng alloy,
ABSTRACT: This Author Certificate introduces a high-resist	lvite allaw
containing 42-50% N1, 40-46% Hn, and 4-18% V.	[ND]
SUB CODE: 11 SUBM DATE: 11May64/ ATD PRESS: 4/18	
요즘 얼마나 하는 사람들은 사람들이 되었다면 하는 사람들이 되었다.	
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그는 소식사회 그는 이 시간 화를 잃으라고 있다. 사람들은 함께 다	
Card 1/1 UDC: 669.245.018.54	

L 09964-67 EWP(e)/EWT(m)/EWP(t)/ETI IJP(c) JD/HW SOURCE CODE: UR/0413/66/000/019/0084/0084 ACC NR AP6035722  R J V : Sorokin, M. N. 36
INVENTOR: Yelyutin, O. P.; Bokshitskiy, I. Ya.; Rogova, I. V.; Sorokin, M. N.
<b>.</b>
ORG: none  TITLE: High-resistivity alloy. Class 40, No. 186694 [announced by the Central TITLE: Research Institute of Ferrous Metallurgy im. Bardina (Tsentral'nyy Scientific Research Institute chernoy metallurgii)]
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Scientific Research Institute of Ferrous Retailed (Scientific Retailed (Scientific Research Institute of Ferrous Retailed (Scientific Reta
SOURCE: Izobreteniya, promyshlennyye obrazesy,
high registivity alloy, titanium containing
SOURCE: Izobreteniya, promyshizany,  TOPIC TAGS: nickel manganese alloy, high resistivity alloy, titanium containing  alloy, cobalt containing alloy
Cartificate introduces a high-resistivity introduces at a
ARSTRACT: This Author Certificate introduces a high-resistivity nickel-manganese- base alloy containing 45—50% nickel, 43—48% manganese, and 2—12% titanium at a base alloy containing 45—50% nickel, 43—48% manganese, and 2—12% titanium and nickel to manganese ratio of 1.0—1.07:1.0. A variant has 5% max titanium and nickel to improve ductility.
nickel to manganese rule to improve ductility.
SUB CODE: 11/ SUBM DATE: 10Aug65/ ATD PRESS: 5105
UDC: 669.018.54: :669.245'74'295
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ACC NR: AP7003259 (N) SOURCE CODE: UR/0207/66/000/006/0103/0104

AUTHOR: Sorokin, M. P. (Perm'); Khlebutin, G. N. (Perm'); Shaydurov, G. F. (Perm')

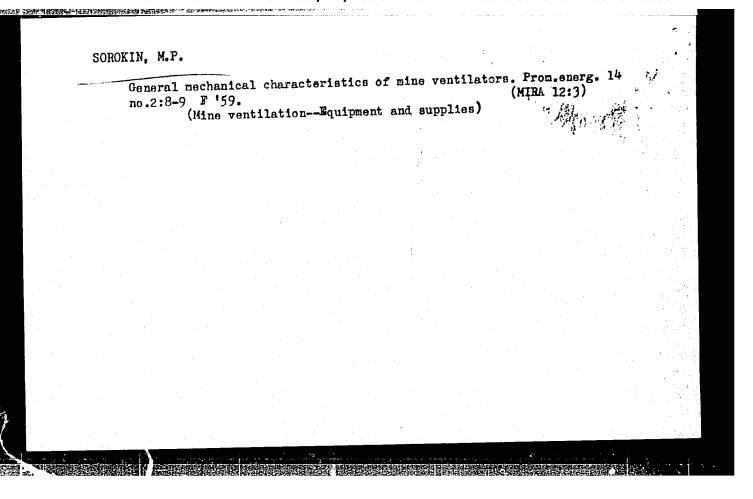
ORG: none

TITLE: Stability of fluid flow between two rotating spherical surfaces

SOURCE: Zhurnal prikladnoy mekhaniki i tekhnicheskoy fiziki, no. 6, 1966, 103-104

TOPIC TAGS: fluid flow, flow stability, boundary layer flow

ABSTRACT: In 1961 it was found that the flow of liquid between two concentric spheres, the inner one in motion and the external one fixed, becomes unstable at the sphere radius ratio  $r_2/r_1 = 2$  with a Reynolds number of about 100 (R =  $r_1^2$   $\omega/v$ , where  $\omega$  is the angular velocity of the rotating sphere and v is kinematic viscosity of the liquid). The solution was sought in powers of the Reynolds number; since the applicability of this method when R is approximately 100 is not obvious, an experiment was set up to test the theory. A special device was constructed with an inner sphere of steel, an external sphere of organic glass, and water or commercially pure glycerine as the fluid. Very precise measurements are possible. Observations showed that there was no qualitative difference in fluid flow at all Reynolds numbers studied from the flow previously found in the second approximation. Considerably less intense motion in the meridian plane is imposed on the circular horizontal motion, i.e., on



SCHOKIN, Mikhail Petrovich; ASTAKHOV, A.V., otv.red.; LOMILINA, L.N., tekhn.red.

[Mine ventilation systems] Shakhtnye ventiliatornye ustanovki.

Moskva, Gos.nauchno-tekhn.izd-vo lit-ry po gornomu delu, 1960.

143 p.

(Mine ventilation—Equipment and supplies)

(Electricity in mining)

SOROKIN, Mikhail Petrovich; LOPATINA, G.G., red.; KARASEV, A.I., tekhn. red.

[Installation, operation, and repair of the electric equipment in metallurgical plants] Montazh, ekspluatatsiia i remont elektrooborudovaniia metallurgicheskikh zavodov. Moskva, Metallurgizdat, 1963. 339 p. (MIRA 16:7) (Metallurgical plants-Electric equipment)

SOROKIN, M.P., inzh.

Measurement of the temperature of electric machinery parts. Vest. elektroprom 34 no.6:13-15 Je '63. (MIRA 16:7)

(Electric machinery)
(Temperature—Measurement)

88634

S/170/61/004/002/013/018 B019/B060

.11.9200

AUTHOR:

Sorokin, M. P.

TITLE:

Experimental Study of the Stability of a Convective Flow

of a Liquid in a Long Vertical Slit

PERIODICAL:

Inzhenerno-fizicheskiy zhurnal, 1961, Vol. 4, No. 2,

, pp. 106-108

TEXT: The experiments described here were performed on a slit whose length 1-to-width d ratio was 1/d=44. One of the plates forming the slit was electrically heatable, while the other could be cooled with water tubes. The power of electric heating decreased linearly from bottom to top. The difference of temperature between the plates was measured with thermocouples. Three to four hours after the system was switched on there appeared a steady state, at which the wall temperature was measured. Three series of experiments were performed, at which the temperature of the cooled slit plate was kept by a thermostat at 11.5, 16.5, and 20.5°C, respectively. Fig. 2 is a graph showing the heated wall temperature

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Experimental Study of the Stability of a Convective Flow of a Liquid in a Long Vertical Slit

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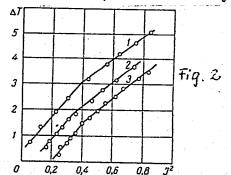
measured in the steady state as a function of the square of the heating current. 1970 (experimental) is obtained for the critical Grashof number, which is near the Grashof number 1910 obtained theoretically by G. Z. Gershuni. A turbulent flow appears above this critical value. There are 2 figures, 1 table, and 7 references: 3 Soviet and 4 British.

ASSOCIATION: Gosudarstvennyy universitet, g. Perm' (State University,

Perm')

SUBMITTED:

August 13, 1960



Card 2/2

25561 S/170/61/004/008/011/016 B125/B201

26.5200

ATTHOR:

Sorokin, M. P.

TITLE:

Convection of a liquid in a cavity under conditions of a

boundary layer

PERIODICAL: Inzhenerno-fizicheskiy zhurnal, v. 4, no. 8, 1961, 107 - 110

TEXT: The steady motion of a liquid in a vertical slit-shaped cavity in the interval from 10<sup>2</sup> to 10<sup>5</sup> of the Grashof numbers has been studied using a device described by G. A. Ostroumov (Svobodnaya Konvektsiya v usloviyakh vnutrenney zadachi (Free convection under conditions of the internal problem), GITTL 1954). The results found by G. F. Shaydurov (ZhTF, XXVIII, no. 4, 1958) concerning convection in a spherical cavity are said to be very interesting. In the middle of a 120°120°40 mm parallelelepiped made of babbitt metal, along ist small axis of symmetry, a 38.6-mm large continuous slit was milled out and covered with window glass. The cavity formed in this way was filled with distilled water. A heat current perpendicular to the slit was then produced using an electric heater and a water-jet cooler. The temperatures on the walls

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Convection of ...

were measured by copper-constantan thermocouples; the temperature field inside the liquid was examined by optical methods. The optics used here differs very little from that suggested by G. A. Ostroumov. The velocity distribution was determined from the motion of light-scattering particles (aluminum powder) suspended in water by photographing the paths of the particles. The character of motion of the liquid in the slit depends largely upon the value of the Grashof number (Gr). For a small Gr, the velocity distribution is in good agreement with the theory applying to an infinite slit with isothermal walls. With rising value of Gr (Gr > 400) a boundary layer appears on the cavity wall, and a little mobile core in the center. The maximum of the velocity profile then approaches the cavity walls, and the temperature practically changes only near them. In this case, the velocity rises in proportion to  ${\rm Gr}^{1/2}$ . When  ${\rm Gr} > 5000$ , a peculiar motion occurs in the middle, as has already been described by G. F. Shaydurov (ZhTF, XXVIII, no. 4, 1958). According to Fig. 1, not only the velocity but also the temperature gradient in the middle is opposed to the corresponding quantity in the boundary layer. The undulatory motion of particles is indicative of a repeated change of the direction of vertical velocity as a function of the horizontal

Card 2/5

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coordinate. The formation of regions with a noticeable motion in a slitshaped cavity can be explained by a vertical temperature gradient. If a
certain parameter mexceeds a definite value, the profile of velocity and
temperature will begin to oscillate. Theory and experiment are in good
agreement; therefore, the appearance of a noticeable motion of the liquid
in the middle is probably due to the vertical temperature gradient. The
formulas

$$v = \frac{1}{4 m^2 M} \left[ \frac{\cos my \sin my}{H} - \frac{\sin my \cosh my}{S} \right], (1)$$

$$T = \frac{1}{2 M} \left[ \frac{\sin my \cosh my}{H} + \frac{\cos my \sinh my}{S} \right] + Az, (2)$$

$$rae$$

$$S = \sin \frac{m}{2} \cosh \frac{m}{2}; \quad H = \cos \frac{m}{2} \sinh \frac{m}{2};$$

$$m = \sqrt[4]{\frac{\text{Gr Pr } A}{4}}; \quad M = \frac{H^2 + S^2}{HS}; \quad A = \frac{\partial T}{\partial z};$$

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used for the abovementioned theoretical calculations are also suited for calculating the stability of convective motion of a liquid in a short slit. Professor G. A. Ostroumov is thanked for having supervised the present work. There are 2 figures and 5 references: 4 Soviet-bloc and 1 non-Soviet-bloc. The reference to the English-language publication reads as follows: Prager S. J. Chem. Phys., XXIII, no. 9, 1955.

ASSOCIATION: Gosudarstvennyy universitet g. Perm' (Perm' State University)

SUBMITTED: December 24, 1960

Card 4/5